

**U. PORTO**

**FMUP** FACULDADE DE MEDICINA  
UNIVERSIDADE DO PORTO

**MESTRADO INTEGRADO EM MEDICINA**

---

2015/2016

João Luís Aragão Rodrigues

Ectropion

março, 2016

FMUP

João Luis Aragão Rodrigues  
Ectropion

**Mestrado Integrado em Medicina**

**Área: Oftalmologia**

**Tipologia: Monografia**

**Trabalho efetuado sob a Orientação de:**  
**Doutora Sara Filipa Teixeira Ribeiro**

**Trabalho organizado de acordo com as normas da revista:**  
**Ophtalmic Plastic and Reconstructive Surgery**

março, 2016

**FMUP**

Eu, João Luis Aragão Rodrigues, abaixo assinado, nº mecanográfico 0608010 42, estudante do 6º ano do Ciclo de Estudos Integrado em Medicina, na Faculdade de Medicina da Universidade do Porto, declaro ter atuado com absoluta integridade na elaboração deste projeto de opção.

Neste sentido, confirmo que **NÃO** incorri em plágio (ato pelo qual um indivíduo, mesmo por omissão, assume a autoria de um determinado trabalho intelectual, ou partes dele). Mais declaro que todas as frases que retirei de trabalhos anteriores pertencentes a outros autores, foram referenciadas, ou redigidas com novas palavras, tendo colocado, neste caso, a citação da fonte bibliográfica.

Faculdade de Medicina da Universidade do Porto, 7 / 01 / 2016

Assinatura conforme cartão de identificação:

João Luis Aragão Rodrigues

**Projecto de Opção do 6º ano – DECLARAÇÃO DE REPRODUÇÃO**

NOME

João Luis Aragão Rodrigues

NÚMERO DE ESTUDANTE

DATA DE CONCLUSÃO

060801042

DESIGNAÇÃO DA ÁREA DO PROJECTO

Oftalmologia

TÍTULO DISSERTAÇÃO/MONOGRAFIA (riscar o que não interessa)

Ectropion

ORIENTADOR

Saua Felipe Teixeira Ribeiro

COORDINADOR (se aplicável)

ASSINALE APENAS UMA DAS OPÇÕES:

É AUTORIZADA A REPRODUÇÃO INTEGRAL DESTA TRABALHO APENAS PARA EFEITOS DE INVESTIGAÇÃO, MEDIANTE DECLARAÇÃO ESCRITA DO INTERESSADO, QUE A TAL SE COMPROMETE.	<input checked="" type="checkbox"/>
É AUTORIZADA A REPRODUÇÃO PARCIAL DESTA TRABALHO (INDICAR, CASO TAL SEJA NECESSÁRIO, Nº MÁXIMO DE PÁGINAS, ILUSTRAÇÕES, GRÁFICOS, ETC.) APENAS PARA EFEITOS DE INVESTIGAÇÃO, MEDIANTE DECLARAÇÃO ESCRITA DO INTERESSADO, QUE A TAL SE COMPROMETE.	<input type="checkbox"/>
DE ACORDO COM A LEGISLAÇÃO EM VIGOR, (INDICAR, CASO TAL SEJA NECESSÁRIO, Nº MÁXIMO DE PÁGINAS, ILUSTRAÇÕES, GRÁFICOS, ETC.) NÃO É PERMITIDA A REPRODUÇÃO DE QUALQUER PARTE DESTA TRABALHO.	<input type="checkbox"/>

Faculdade de Medicina da Universidade do Porto, 5 / 04 / 2016

Assinatura conforme cartão de identificação:

João Luis Aragão Rodrigues

À minha mãe, ao meu pai e à minha irmã.

# ECTROPION

Sara F T Ribeiro, MD, PhD<sup>1,3</sup>, João L A Rodrigues<sup>2</sup>, Maria Shekhovtsova, MD<sup>2</sup>,  
Antonio A V Cruz, MD, PhD<sup>3</sup>

<sup>1</sup> Department of Ophthalmology, Hospital de Braga, School of Health Sciences,  
University of Minho, Braga, Portugal.

<sup>2</sup> School of Medicine of Porto, University of Porto, Porto, Portugal.

<sup>3</sup> Department of Ophthalmology, Otorhinolaryngology and Head and Neck Surgery,  
Hospital das Clínicas-Campus, School of Medicine of Ribeirão Preto, University of  
São Paulo, Ribeirão Preto, São Paulo, Brazil.

**Runing head:** Ectropion.

**Corresponding author:** Sara Filipa Teixeira Ribeiro

Postal address: Department of Ophthalmology, Hospital de Braga, School of Health  
Sciences, University of Minho, Rua Sete Fontes – São Vítor 4710-243 Braga,  
Portugal.

Email: sara\_ribeiro25@hotmail.com

Telephone: 00351-919589546

21    **Précis**

22    Ectropion is an abnormal eversion of the eyelid margin. The understanding of the  
23    underlying pathophysiological mechanism is the key to a successful surgical  
24    treatment.

## ABSTRACT

**Purpose:** Ectropion is one of the most common eyelid malpositions. The treatment of ectropion is still a subject of controversy. The authors reviewed the literature about ectropion, including definition, diagnosis, pathophysiological mechanisms, classification and surgical treatments.

**Methods:** A literature search was performed on the MEDLINE database using the keywords ectropion, eyelid malposition, cicatricial ectropion, congenital ectropion, involutional ectropion, mechanical ectropion, tarsal strip; lazy-T; canthoplasty, canthopexy, skin flaps, skin grafts. Only articles in english were included.

**Results:** There is no consensus about the best surgical treatment for this eyelid malposition, however, we know that the identification of the anatomic abnormality associated is the most important point to chose the correct surgical treatment. The literature on the treatment of ectropion mainly includes descriptions of surgical techniques without objective measurements of the results, and uncontrolled studies.

**Conclusions:** The scientific literature on ectropion is vast, however there is a need for quantitative studies on the effects of ectropion correction using different surgical techniques.



Ectropion is an eversion (outward turning) of the eyelid margin. This lid malposition may affect the lower or upper lid and leads to cosmetic and functional deficits, such as conjunctival hyperemia, corneal exposure, eye irritation, epiphora, chronic conjunctivitis, and in rare cases visual loss. It can be medial or lacrimal, lateral or complete (involving the entire eyelid).<sup>1,2</sup>

## **METHODS OF LITERATURE SEARCH**

A literature search was performed on the MEDLINE database using the keywords ectropion, eyelid malposition, cicatricial ectropion, congenital ectropion, involutional ectropion, mechanical ectropion, tarsal strip; lazy-T; canthoplasty, canthopexy, skin flaps, skin grafts. Only articles in english were included.

The research adhered to the tenets of the Declaration of Helsinki and was approved by the Ethics Committee of the Hospital de Braga, Braga, Portugal. An informed consent was obtained from patients whose photos were used in this article.

## **CLASSIFICATION OF ECTROPION**

Ectropion is usually classified as congenital, involutional, cicatricial, mechanical and paralytic.<sup>1,2</sup> These categories are not absolute and some cases fit in more than one type. For instance, a four-lid ectropion in a newborn with lamellar ichthyosis can be considered as a congenital or a cicatricial ectropion. Similarly, the ectropion of a child born with an orbital cyst, which mechanically pushes the lid margin, may be seen as a congenital mechanical ectropion, and congenital facial palsies may cause a congenital paralytic ectropion.

From a therapeutic perspective, rather than focusing on the clinical categories it is more useful to look at the underlying pathophysiological mechanism (Table 1) because the correct identification of the anatomic abnormality associated with the lid margin malposition is the key to a successful surgical procedure.

## **CONGENITAL ECTROPION**

The designation “congenital ectropion” is used to name a variety of conditions with distinct pathophysiological mechanisms. A typical example is the so-called congenital upper eyelid eversion. This rare lid abnormality, usually present at birth, is characterized by eversion of both upper lid margins with prolapsed chemotic conjunctiva. It is typically symmetrical but some degree of asymmetry is not uncommon. Several factors have been implicated in its pathophysiology including trauma at birth, orbicularis hypotonia, posterior lamella vertical elongation and failure of the orbital septum to fuse with the levator aponeurosis with adipose tissue interposition. Most cases are not associated with ocular or general abnormalities. However, the incidence appears to be higher in Down syndrome. This peculiar form of upper lid ectropion responds well to conservative treatment such as topical ointments and lubricants, patching or temporary tarsorrhaphy.<sup>3-6</sup>

A totally different situation is represented by newborns with severe forms of lamellar ichthyosis. In these patients eyelid eversion is due to severe skin contraction. All four lids may undergo a severe form of cicatricial ectropion that might require early skin grafting in order to prevent corneal ulceration and eye perforation. Shortening of the anterior lamella is also implicated in other syndromes associated with congenital ectropion such as blepharophimosis (lower lid laterally)<sup>7</sup>,

blepharochelodontic syndrome (lower eyelids)<sup>8-10</sup> and Down syndrome (upper and lower lids)<sup>11</sup>, as well a variety of sporadic congenital anomalies including craniofacial clefting.<sup>12</sup>

## **INVOLUTIONAL ECTROPION**

Involutional ectropion is the most frequent form of lower eyelid eversion, commonly found in aged patients. The Blue Mountain Eye Study assessed the prevalence and associations of ectropion in a large cohort of residents of Sydney, Australia (3654 people aged 49-97 years). Ectropion prevalence was higher in men (5.1%) than women (3.0%). The prevalence of lower eyelid ectropion increased with age, reaching 16.7% among patients above 80 years old.<sup>13</sup> Damasceno studied a Brazilian population of 24 565 elderly people and found similar results, with a prevalence of 17.7% in subjects aged 80 years or more.<sup>14</sup>

Several factors have been implicated in the pathogenesis of this eyelid deformity, all related to an abnormal laxity of the lid support system including tarsal and orbital septum atrophy, thinning of the skin and subcutaneous tissues, elongation of the tarsus and pretarsal orbicularis, medial and lateral canthal tendon laxity and dehiscence, and elongation or desinsertion of the lower lid retractors.<sup>2</sup> Heimmell suggested that when these predisposing factors are present, the key to final lower eyelid position is the globe axial projection, with relatively exophthalmic eyes being more likely to develop tarsal ectropion.<sup>15</sup>

Histopathological studies have confirmed the role of age-related changes in the tarsal plate, inferior retractors, orbicularis oculi and lateral canthal tendon in the

lids with involutional ectropion.<sup>16,17</sup> The affected lid shows the presence of collagen degeneration and elastosis of the tarsal plate and canthal tendons, an increased amount of adipose tissue in the distal tarsus, and focal degeneration, fibrosis, and elastosis of the pretarsal orbicularis.<sup>18-20</sup> In chronic sun exposure, actinic damage on the anterior lamella contributes as an additional factor to lower eyelid eversion in patients with involutional ectropion.<sup>21</sup>

The evaluation of a patient with involutional ectropion should start with the inspection of the lid margin and the position of the puncta since one the first manifestations of lower eyelid ectropion is epiphora secondary to lacrimal punctum eversion. The patency of the puncta and lacrimal drainage system should always be examined.

The lower eyelid laxity can be detected using the “pinch test” and the “snap back test”. The result is abnormal if the lid can be distended more than 6 mm from the globe or does not briskly return to its natural position. The lateral canthus should form an acute angle and a rounded shape is indicative of lateral tendon elongation. If during traction of the lower lid there is lateral displacement of the lacrimal punctum towards the limbus, there is medial tendon laxity.<sup>2, 22</sup>

The condition of the lower lid skin is assessed by pulling the lower lid margin upwards. With traction the lid margin should reach a point at least 2 mm above the limbus. A relatively immobile margin indicates vertical shortening of the anterior lamella. This cicatricial component of the involutional ectropion is a common finding when the ectropion is present for a long period of time or can be the result of mild actinic changes of the skin.<sup>2, 22</sup>

## CICATRICIAL ECTROPION

Cicatricial ectropion is caused by anterior lamella shortening. Depending on the etiologic mechanism, cicatricial entropion can affect the upper lids, the lower lids or both (as observed in some cases of ichthyosis). It usually involves secondary eyelid scars resulting from trauma<sup>23, 24</sup>, burns<sup>25-27</sup> or from a large contingent of skin diseases such as ichthyoses<sup>28, 29</sup>, discoid lupus erythematosus<sup>30</sup>, inherited epidermolysis bullosa<sup>31</sup>, generalized eruptive keratoacanthoma<sup>32-34</sup>, cutaneous leishmaniasis<sup>35</sup>, pityriasis rubra pilaris<sup>36</sup>, and pyoderma gangrenosum<sup>37, 38</sup>.

Postoperative complications of eyelid tumors<sup>39, 40</sup>, blepharoplasty<sup>41-44</sup> and skin resurfacing<sup>45, 46</sup> are also common causes of lower lid cicatricial ectropion. Other sources of skin damage resulting in ectropion are radiotherapy<sup>47, 48</sup> and the use of drugs such as docetaxel<sup>49</sup>, fluorouracil<sup>50-52</sup>, anthrax<sup>53</sup>, prostaglandins<sup>54, 55</sup>, brimonidine<sup>55</sup>, betaxolol<sup>55</sup>, dorzolamide<sup>55</sup>, timolol<sup>55</sup>, and iopidine<sup>56</sup>.

## PARALYTIC ECTROPION

Paralytic ectropion is caused by the lack of normal innervation of the orbicularis muscle. Failure of normal lid closure with lower lid laxity and ectropion, upper eyelid retraction and brow ptosis are the clinical signs. The causes of seventh nerve palsy are myriad, but can be broadly divided, in the order of frequency, into idiopathic (Bell's palsy), traumatic (birth canal trauma or forceps delivery, surgical trauma, facial and temporal bone fractures), infectious (herpes zoster, Lyme disease, HIV infection, polio, mumps, cytomegalovirus, mononucleosis, leprosy, cat scratch fever), and neoplastic (acoustic neuroma, cerebellopontine angle tumors).<sup>57</sup>

In paralytic ectropion due to surgical trauma, the timing for facial reanimation surgery is under debate. Watts *et al* advocate that rehabilitation surgery with gold weight implantation and lateral tarsal strip should be performed immediately, at the time of facial nerve sacrifice.<sup>58</sup>

## **MECHANICAL ECTROPION**

Mechanical ectropion is caused by eyelid tumors that evert the lower lid or inflammatory disorders that cause orbicularis spasm. Large tumors or cysts near the lid margin, acute proptosis with chemosis, eyelid and periocular edema, significant herniated orbital fat and traction on the lower eyelid skin from spectacles can mechanically cause ectropion. The treatment is directed at the cause.<sup>2, 59-62</sup>

## **TREATMENT**

Lower or upper eyelid ectropions are usually managed with surgery. As mentioned before, the key for a successful surgical procedure is the correct preoperative identification of the underlying etiologic factor. Depending on the mechanism provoking the lid margin rotation, a variety of procedures are used to stabilize the lid margin. For instance, lower eyelid horizontal laxity with normal canthal tendon tonus can be addressed by pentagonal full thickness eyelid resection. If the lateral canthal tendon is abnormally lax tarsal strip procedures are indicated.<sup>1, 2, 63, 64</sup> This useful procedure is a variant of the old Bick's lateral resection<sup>65</sup> and consists of a lateral canthotomy and cantholysis; excision of skin and conjunctiva, leaving a free strip of tarsus; fixation of the tarsal strip to the periosteum of the lateral

orbital wall; and reconstruction of the lateral canthus to create the appropriate height and tension of the lower eyelid.<sup>63, 66</sup> Usually monofilament non-absorbable sutures are used<sup>2, 64, 67, 68</sup>, but long-acting absorbable sutures will also work without recurrence of horizontal laxity.<sup>66, 69, 70</sup>

The repair of medial tendon laxity is more complicated than that of its lateral counterpart because of the intimate relation of the tendon with the canaliculi. The high rate of postoperative lacrimal problems following surgery on the medial canthus leads some surgeons to delay correction of the medial canthal tendon until laxity is advanced. Medial canthal tendon correction has been attempted by resection or plication of the tendon medial to the lower punctum, anchoring the tissue to either the anterior periosteum or posterior lacrimal crest.<sup>71-75</sup>

Lacrimal ectropion without horizontal laxity is classically treated with an excision of a diamond of conjunctiva and retractors below the punctum. If a horizontal laxity is present, a full-thickness wedge excision horizontally tight of the lid margin can be associated with the pentagon excision creating a lazy-T procedure. The wedge excision can be displaced laterally or be replaced by a tarsal strip.<sup>76, 77</sup>

In both procedures the role of lower lid retractor plication has been emphasized. Advanced tarsal ectropions require a large excision of conjunctiva and retractors by a posterior approach combined with everting sutures and horizontal tightening.<sup>78-80</sup>

Since cicatricial ectropion is due to an anterior lamella deficiency, various surgical techniques can be used to lengthen the underlying deformity. Z-plasties can be used to manage linear scars.<sup>1, 2, 81</sup> Large shortages of skin require local flaps<sup>26, 82-85</sup> or skin grafts<sup>27, 48, 86</sup>. There are a large variety of flaps that can be used to

208 correct lower and upper ectropions depending on the location and extension of the  
209 cicatricial process. Free skin grafts may be obtained from the upper lid, retroauricular  
210 region, supraclavicular area or inner aspect of the upper arm.<sup>1, 2, 81</sup>

211 In severe lower eyelid ectropion, there is often inadequate muscular support  
212 for the pretarsal lower eyelid. When the pretarsal orbicularis muscle is damaged, a  
213 fascia lata sling is an option for supporting the lower lid.<sup>84</sup>

214

## 215 **CONCLUSION**

216 A successful functional and cosmetic correction of ectropion depends on the  
217 understanding that ectropion is a group of eyelid malpositions that may have  
218 different etiological factors and may affect lateral, central or medial portion of the lids,  
219 alone or in combination. Surgery techniques must address the anatomic  
220 abnormalities responsible for the ectropion. The choice of the right procedure leads  
221 to a successful outcome for the patient which is the main goal.



222   **REFERENCES**

- 223   1. Hintschich C. Correction of entropion and ectropion. *Dev Ophthalmol* 2008;41:85-102.
- 224   2. Bedran EG, Pereira MV, Bernardes TF. Ectropion. *Semin Ophthalmol* 2010;25:59-65.
- 225   3. Dohvoma VA, Nchifor A, Ngwanou AN, et al. Conservative management in congenital
- 226   bilateral upper eyelid eversion. *Case Rep Ophthalmol Med* 2015;2015:389289.
- 227   4. Krishnappa NC, Deb AK, Poddar C. Congenital total eversion of upper eyelids in a newborn
- 228   with Down's syndrome. *Oman J Ophthalmol* 2014;7:98-9.
- 229   5. Ibraheem WA. Bilateral congenital upper eyelid eversion: the clinical course and outcome
- 230   of conservative management. *Pan Afr Med J* 2014;17:215.
- 231   6. Fasina O. Management of bilateral congenital upper eyelid eversion with severe
- 232   chemosis. *J Ophthalmic Vis Res* 2013;8:175-8.
- 233   7. Sandramouli S, Betharia SM. Blepharophimosis syndrome: an atypical case. *Eye (Lond)*
- 234   1994;8:482-4.
- 235   8. Gorlin RJ, Wiedemann HR. Blepharo-cheilo-dontic (BCD) syndrome. *Acta Ophthalmol*
- 236   *Scand Suppl* 1996;219:22.
- 237   9. Guion-Almeida ML, Rodini ES, Kokitsu-Nakata NM, Bologna-Amantini D. Blepharo-Cheilo-
- 238   Dontic (BCD) syndrome: report on four new patients. *Am J Med Genet* 1998;76:133-6.
- 239   10. Yen MT, Lucci LM, Anderson RL. Management of eyelid anomalies associated with
- 240   Blepharo-cheilo-dontic syndrome. *Am J Ophthalmol* 2001;132:279-80.
- 241   11. Sellar PW, Bryars JH, Archer DB. Late presentation of congenital ectropion of the eyelids
- 242   in a child with Down syndrome: a case report and review of the literature. *J Pediatr*
- 243   *Ophthalmol Strabismus* 1992;29:64-7.
- 244   12. Iida A, Narai S, Takagi R, et al. Blepharo-cheilo-dontic (BCD) syndrome: case report. *Cleft*
- 245   *Palate Craniofac J* 2006;43:237-43.

246 13. Mitchell P, Hinchcliffe P, Wang JJ, et al. Prevalence and associations with ectropion in an  
247 older population: the Blue Mountains Eye Study. *Clin Experiment Ophthalmol* 2001;29:108-  
248 10.

249 14. Damasceno RW, Osaki MH, Dantas PE, Belfort R, Jr. Involutional entropion and ectropion  
250 of the lower eyelid: prevalence and associated risk factors in the elderly population. *Ophthal*  
251 *Plast Reconstr Surg* 2011;27:317-20.

252 15. Heimmell MR, Enzer YR, Hofmann RJ. Entropion-ectropion: the influence of axial globe  
253 projection on lower eyelid malposition. *Ophthal Plast Reconstr Surg* 2009;25:7-9.

254 16. Chua J, Choo CT, Seah LL, et al. A 5-year retrospective review of Asian ectropion: how  
255 does it compare to ectropion amongst non-Asians? *Ann Acad Med Singapore* 2011;40:84-9.

256 17. Bashour M, Harvey J. Causes of involutional ectropion and entropion--age-related tarsal  
257 changes are the key. *Ophthal Plast Reconstr Surg* 2000;16:131-41.

258 18. Kocaoglu FA, Katircioglu YA, Tok OY, et al. The histopathology of involutional ectropion  
259 and entropion. *Can J Ophthalmol* 2009;44:677-9.

260 19. Damasceno RW, Osaki MH, Dantas PE, Belfort R, Jr. Involutional ectropion and  
261 entropion: clinicopathologic correlation between horizontal eyelid laxity and eyelid  
262 extracellular matrix. *Ophthal Plast Reconstr Surg* 2011;27:321-6.

263 20. Damasceno RW, Heindl LM, Hofmann-Rummelt C, et al. Pathogenesis of involutional  
264 ectropion and entropion: the involvement of matrix metalloproteinases in elastic fiber  
265 degradation. *Orbit* 2011;30:132-9.

266 21. Marshall JA, Valenzuela AA, Strutton GM, Sullivan TJ. Anterior lamella actinic changes as  
267 a factor in involutional eyelid malposition. *Ophthal Plast Reconstr Surg* 2006;22:192-4.

268 22. Frueh BR, Schoengarth LD. Evaluation and treatment of the patient with ectropion.  
269 *Ophthalmology* 1982;89:1049-54.

270 23. Desciak EB, Eliezri YD. Surgical Pearl: Temporary suspension suture (Frost suture) to help  
271 prevent ectropion after infraorbital reconstruction. *J Am Acad Dermatol* 2003;49:1107-8.

272 24. Salgarelli AC, Bellini P, Multinu A, Landini B, Consolo U. Tarsal strip technique for  
273 correction of malposition of the lower eyelid after treatment of orbital trauma. *Br J Oral*  
274 *Maxillofac Surg* 2009;47:298-301.

275 25. Astori IP, Muller MJ, Pegg SP. Cicatricial, postburn ectropion and exposure keratitis.  
276 *Burns* 1998;24:64-7.

277 26. Kostakoglu N, Ozcan G. Orbicularis oculi myocutaneous flap in reconstruction of  
278 postburn lower eyelid ectropion. *Burns* 1999;25:553-7.

279 27. Mandrekas AD, Zambacos GJ, Anastasopoulos A. Treatment of bilateral severe eyelid  
280 burns with skin grafts: an odyssey. *Burns* 2002;28:80-6.

281 28. Cruz AA, Menezes FA, Chaves R, et al. Eyelid abnormalities in lamellar ichthyoses.  
282 *Ophthalmology* 2000;107:1895-8.

283 29. Craiglow BG, Choate KA, Milstone LM. Topical tazarotene for the treatment of ectropion  
284 in ichthyosis. *JAMA Dermatol* 2013;149:598-600.

285 30. Kopsachilis N, Tsaousis KT, Tourtas T, Tsinopoulos IT. Severe chronic blepharitis and  
286 scarring ectropion associated with discoid lupus erythematosus. *Clin Exp Optom*  
287 2013;96:124-5.

288 31. Fine JD, Johnson LB, Weiner M, et al. Eye involvement in inherited epidermolysis bullosa:  
289 experience of the National Epidermolysis Bullosa Registry. *Am J Ophthalmol* 2004;138:254-  
290 62.

291 32. Consigli JE, Gonzalez ME, Morsino R, et al. Generalized eruptive keratoacanthoma  
292 (Grzybowski variant). *Br J Dermatol* 2000;142:800-3.

293 33. Oakley A, Ng S. Grzybowski's generalized eruptive keratoacanthoma: remission with  
294 cyclophosphamide. *Australas J Dermatol* 2005;46:118-23.

295 34. Anzalone CL, Cohen PR. Generalized eruptive keratoacanthomas of Grzybowski. *Int J*  
296 *Dermatol* 2014;53:131-6.

297 35. Chaudhry IA, Hylton C, DesMarchais B. Bilateral ptosis and lower eyelid ectropion  
298 secondary to cutaneous leishmaniasis. *Arch Ophthalmol* 1998;116:1244-5.

299 36. Durairaj VD, Horsley MB. Resolution of pityriasis rubra pilaris-induced cicatricial  
300 ectropion with systemic low-dose methotrexate. *Am J Ophthalmol* 2007;143:709-10.

301 37. Procianoy F, Barbato MT, Osowski LE, Bocaccio FJ, Bakos L. Cicatricial ectropion  
302 correction in a patient with pyoderma gangrenosum: case report. *Arq Bras Oftalmol*  
303 2009;72:384-6.

304 38. Thampy RS, Al-Niaimi F, Lyon C, Duff CG, Leatherbarrow B. Management of cicatricial  
305 ectropion secondary to pyoderma gangrenosum. *Orbit* 2014;33:129-31.

306 39. Rubin P, Mykula R, Griffiths RW. Ectropion following excision of lower eyelid tumours  
307 and full thickness skin graft repair. *Br J Plast Surg* 2005;58:353-60.

308 40. Salgarelli AC, Francomano M, Magnoni C, Bellini P. Cicatricial iatrogenic lower eyelid  
309 malposition in skin cancer surgery: results of a combined approach. *J Craniomaxillofac Surg*  
310 2012;40:579-83.

311 41. Botti G. Blepharoplasty: A classification of selected techniques in the treatment and  
312 prevention of lower lid margin distortions. *Aesthetic Plast Surg* 1998;22:341-8.

313 42. Morax S, Touitou V. Complications of blepharoplasty. *Orbit* 2006;25:303-18.

314 43. Patrocinio TG, Loredó BA, Arevalo CE, et al. Complications in blepharoplasty: how to  
315 avoid and manage them. *Braz J Otorhinolaryngol* 2011;77:322-7.

316 44. Mack WP. Complications in periocular rejuvenation. *Facial Plast Surg Clin North Am*  
317 2010;18:435-56.

318 45. Demas PN, Bridenstine JB. Diagnosis and treatment of postoperative complications after  
319 skin resurfacing. *J Oral Maxillofac Surg* 1999;57:837-41.

320 46. Fife DJ, Fitzpatrick RE, Zachary CB. Complications of fractional CO2 laser resurfacing: four  
321 cases. *Lasers Surg Med* 2009;41:179-84.

322 47. Tarallo M, Rizzo MI, Monarca C, et al. Optimal care for eyelid contraction after  
323 radiotherapy: case report and literature review. *J Oral Maxillofac Surg* 2012;70:2459-65.

324 48. Kim HJ, Hayek B, Nasser Q, Esmaeli B. Viability of full-thickness skin grafts used for  
325 correction of cicatricial ectropion of lower eyelid in previously irradiated field in the  
326 periocular region. *Head Neck* 2013;35:103-8.

327 49. Kaya AO, Buyukberber S, Coskun U, et al. Acute erythema and edematous skin reaction  
328 and ectropion following docetaxel in a patient with non-small cell lung cancer. *Cutan Ocul*  
329 *Toxicol* 2008;27:327-31.

330 50. Lewis JE. Temporary ectropion due to topical fluorouracil. *Int J Dermatol* 1997;36:79.

331 51. Obi EE, McDonald A, Kemp E. A bilateral cicatricial ectropion and bilateral upper lid  
332 shortening caused by 5-fluorouracil toxicity in a patient with dihydropyrimidine  
333 dehydrogenase deficiency. *Cutan Ocul Toxicol* 2011;30:157-9.

334 52. Nikkhah D, Abood A, Watt D. Cicatricial ectropion: a complication of topical 5-  
335 fluorouracil. *J Plast Reconstr Aesthet Surg* 2012;65:e9-10.

336 53. Soysal HG, Kiratli H, Recep OF. Anthrax as the cause of preseptal cellulitis and cicatricial  
337 ectropion. *Acta Ophthalmol Scand* 2001;79:208-9.

338 54. Altieri M, Ferrari E. Do prostaglandin analogs affect eyelid position and motility? *J Ocul*  
339 *Pharmacol Ther* 2011;27:511-7.

340 55. Hegde V, Robinson R, Dean F, Mulvihill HA, Ahluwalia H. Drug-induced ectropion: what is  
341 best practice? *Ophthalmology* 2007;114:362-6.

342 56. Britt MT, Burnstine MA. Iopidine allergy causing lower eyelid ectropion progressing to  
343 cicatricial entropion. *Br J Ophthalmol* 1999;83:992-3.

344 57. Mavrikakis I. Facial nerve palsy: anatomy, etiology, evaluation, and management. *Orbit*  
345 2008;27:466-74.

346 58. Watts TL, Chard R, Weber SM, Wax MK. Immediate eye rehabilitation at the time of  
347 facial nerve sacrifice. *Otolaryngol Head Neck Surg* 2011;144:353-6.

348 59. Ashfaq I, Kyprianou I, Ahluwalia H. A large kissing (divided) naevus presenting with  
349 complete mechanical ptosis and lower lid ectropion. *J Plast Reconstr Aesthet Surg*  
350 2009;62:e87-8.

351 60. Kampp JT, Kouba DJ, Fincher EF, Moy RL. Basal cell carcinoma masquerading as the  
352 chronic ectropion of lamellar ichthyosis. *Dermatol Surg* 2008;34:963-7.

353 61. Baker HE, Berry-Brincat A, Zaki I, Cheung D. Three different consecutive manifestations  
354 of morphoeic BCC in the same patient: presenting first as ectropion, then entropion, and  
355 finally medial canthal dystopia with epicanthus inversus. *Orbit* 2008;27:183-5.

356 62. Evans M, Chang E, Yu DL, Rao NA. Granular cell tumour: a rare caruncle lesion. *Br J*  
357 *Ophthalmol* 2006;90:246-7.

358 63. Anderson RL, Gordy DD. The tarsal strip procedure. *Arch Ophthalmol* 1979;97:2192-6.

359 64. Della Rocca DA. The lateral tarsal strip: illustrated pearls. *Facial Plast Surg* 2007;23:200-  
360 2.

361 65. Bick MW. Surgical management of orbital tarsal disparity. *Arch Ophthalmol* 1966;75:386-  
362 9.

363 66. Jordan DR, Anderson RL. The lateral tarsal strip revisited. The enhanced tarsal strip. *Arch*  
364 *Ophthalmol* 1989;107:604-6.

365 67. Kam KY, Cole CJ, Bunce C, Watson MP, Kamal D, Olver JM. The lateral tarsal strip in  
366 ectropion surgery: is it effective when performed in isolation? *Eye (Lond)* 2012;26:827-32.

367 68. Tenzel RR, Buffam FV, Miller GR. The use of the "lateral canthal sling" in ectropion  
368 repair. *Can J Ophthalmol* 1977;12:199-202.

369 69. Hsuan J, Selva D. The use of a polyglactin suture in the lateral tarsal strip procedure. *Am*  
370 *J Ophthalmol* 2004;138:588-91.

371 70. Meyer DR. The use of a polyglactin suture in the lateral tarsal strip procedure. *Am J*  
372 *Ophthalmol* 2005;139:758-9.

373 71. O'Donnell BA, Anderson RL, Collin JR, et al. Repair of the lax medial canthal tendon. *Br J*  
374 *Ophthalmol* 2003;87:220-4.

375 72. Moe KS, Kao CH. Precaruncular medial canthopexy. *Arch Facial Plast Surg* 2005;7:244-  
376 50.

377 73. Sullivan TJ, Collin JR. Medial canthal resection: an effective long-term cure for medial  
378 ectropion. *Br J Ophthalmol* 1991;75:288-91.

379 74. Francis IC, Wilcsek GA, Sharma S, Coroneo MT. Transcaruncular medial orbitotomy for  
380 stabilization of the posterior limb of the medial canthal tendon. *Clin Experiment Ophthalmol*  
381 2001;29:85-9.

382 75. Edelstein JP, Dryden RM. Medial palpebral tendon repair for medial ectropion of the  
383 lower eyelid. *Ophthalm Plast Reconstr Surg* 1990;6:28-37.

384 76. Manners RM. Surgical repair of medial ectropion. *Eye (Lond)* 1995;9:365-7.

385 77. Smith B. The "lazy-T" correction of ectropion of the lower punctum. *Arch Ophthalmol*  
386 1976;94:1149-50.

387 78. Ferguson AW, Chadha V, Kearns PP. The not-so-lazy-T: a modification of medial  
388 ectropion repair. *Surgeon* 2006;4:87-9.

389 79. O'Donnell B. Age-related medial ectropion of the lower eyelid. *Aust N Z J Ophthalmol*  
390 1994;22:183-6.

391 80. Fong KC, Mavrikakis I, Sagili S, Malhotra R. Correction of involutional lower eyelid medial  
392 ectropion with transconjunctival approach retractor plication and lateral tarsal strip. *Acta*  
393 *Ophthalmol Scand* 2006;84:246-9.

394 81. Miletic D, Elabjer BK, Bosnar D, Busic M. Our approach to operative treatment of lower  
395 lid ectropion. *Acta Clin Croat* 2010;49:283-7.

396 82. Xue CY, Dai HY, Li L, et al. Reconstruction of lower eyelid retraction or ectropion using a  
397 paranasal flap. *Aesthetic Plast Surg* 2012;36:611-7.

398 83. Xu JH, Tan WQ, Yao JM. Bipedicle orbicularis oculi flap in the reconstruction of the lower  
399 eyelid ectropion. *Aesthetic Plast Surg* 2007;31:161-6.

400 84. Qian JG, Wang XJ, Wu Y. Severe cicatrical ectropion: repair with a large advancement  
401 flap and autologous fascia sling. *J Plast Reconstr Aesthet Surg* 2006;59:878-81.

402 85. Manku K, Leong JK, Ghabrial R. Cicatricial ectropion: repair with myocutaneous flaps and  
403 canthopexy. *Clin Experiment Ophthalmol* 2006;34:677-81.

404 86. Lau CK, Huang S, Cormack G. Minimising the risk of ectropion when full thickness skin  
405 grafting lower eyelid defects. *J Plast Reconstr Aesthet Surg* 2008;61:1562-4.



**Table 1.** Type of ectropion, affected lids and pathophysiologic mechanism.

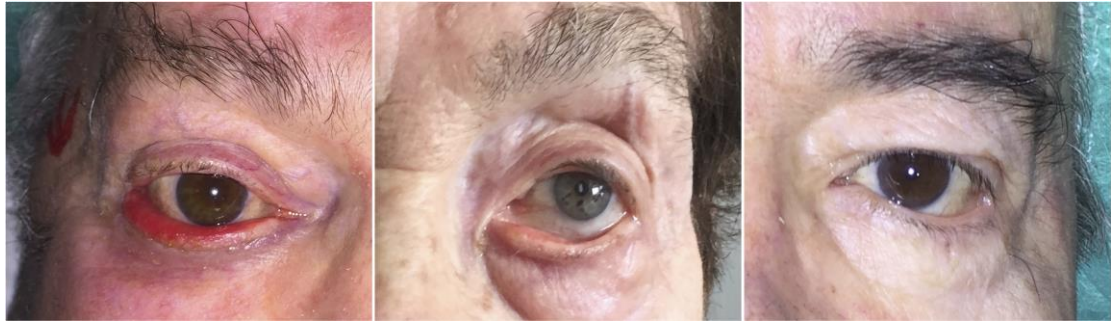
Type of Ectropion	Affected Lid (s)	Pathophysiologic Mechanism
Congenital	UL, LL	Tarsus anomaly (absence or atrophy)
		Severe eyelid edema after birth trauma
		Eyelid skin retraction (cicatricial)
		Microphthalmus with orbital cyst (mechanical)
Involutional	LL	Horizontal lid laxity
		Lateral and/or medial canthal tendon laxity
		Tarsal and orbital septum atrophy
		Thinning of skin and subcutaneous tissues
		Senile enophthalmia
		Dehiscence, elongation or desinsertion of the lower lid retractors
Cicatricial	UL, LL	Shortening of the anterior lamella of the lid
Paralytic	LL	Orbicularis muscle atony
Mechanical	LL	Lesions or inflammatory disorders which cause the lid margin to roll out

UL, upper lid. LL, lower lid.

**Figure 1.** Lamellar ichthyosis. Congenital cicatricial ectropion of the lower eyelids.



**Figure 2.** Involutional ectropion: A) total; B) medial; C) lateral.



**Figure 2.** Cicatricial ectropion caused by an excision of basal cell carcinoma of the lower eyelid.

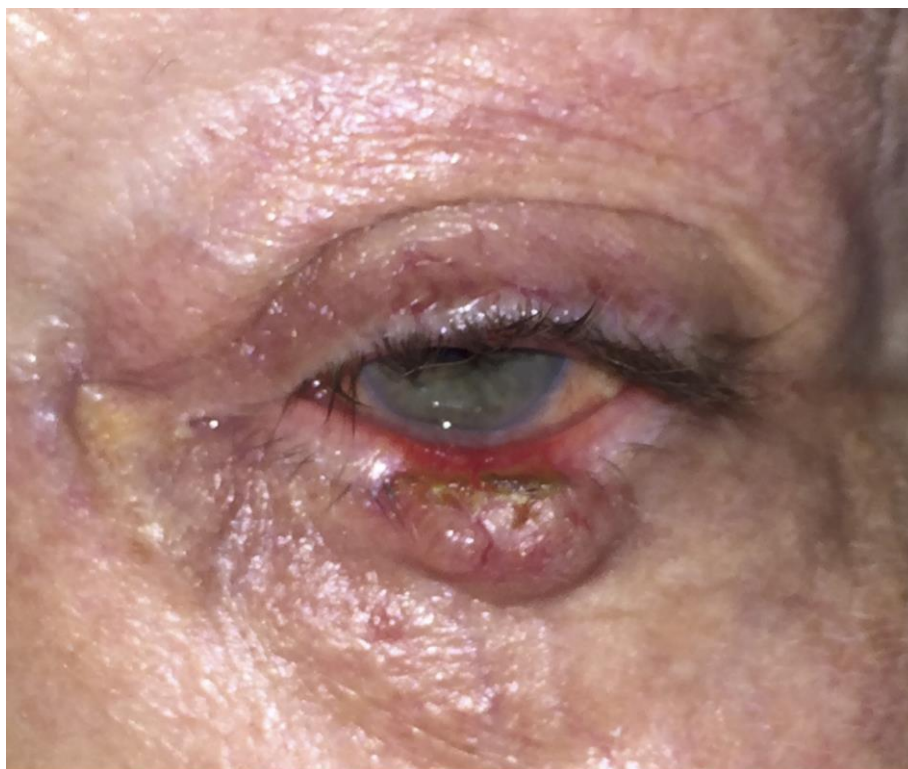




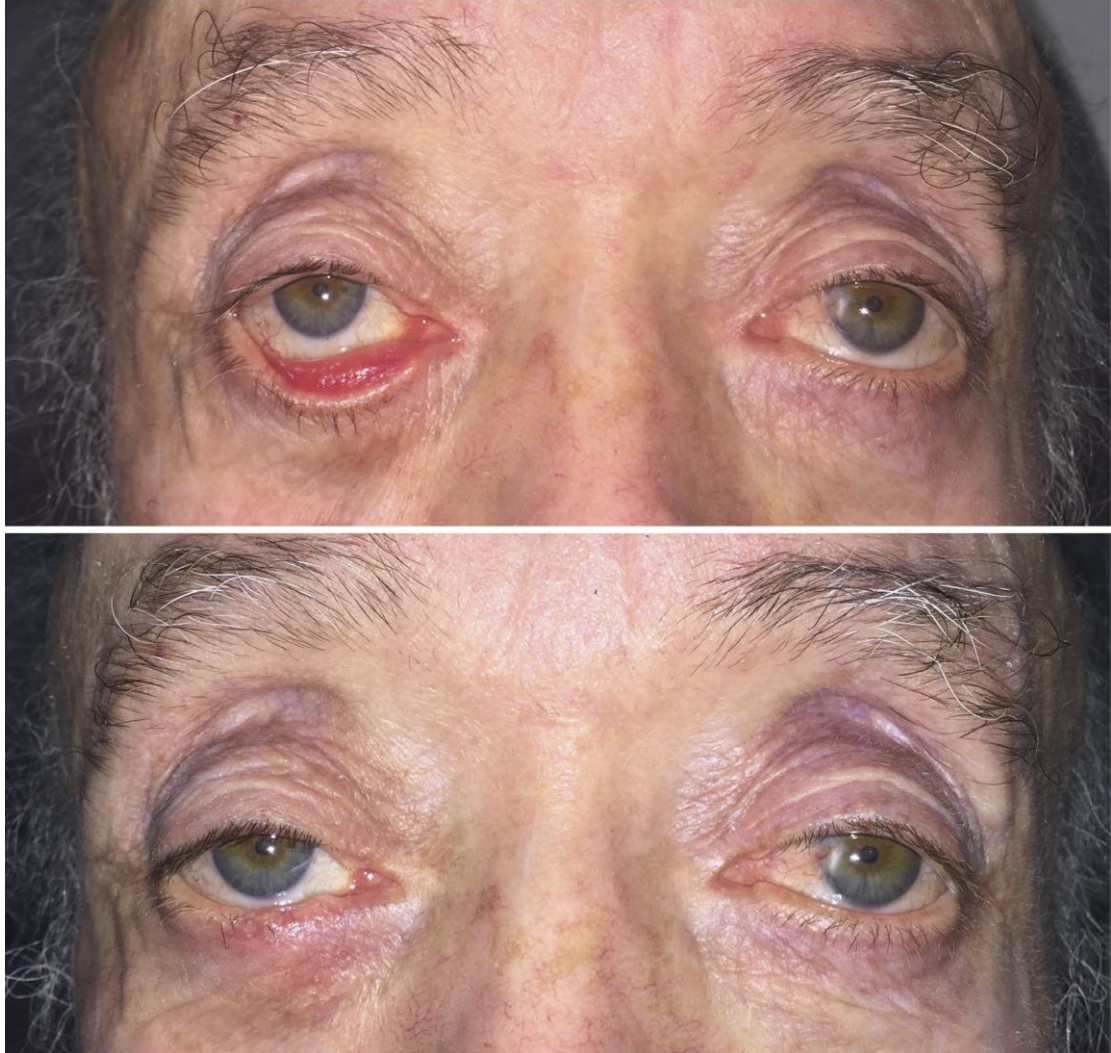
**Figure 4.** Left lower lid paralytic ectropion. Note lower lid laxity and ectropion, brow ptosis and absence of forehead wrinkles.



**Figure 5.** Mechanical ectropion caused by a basal cell carcinoma.

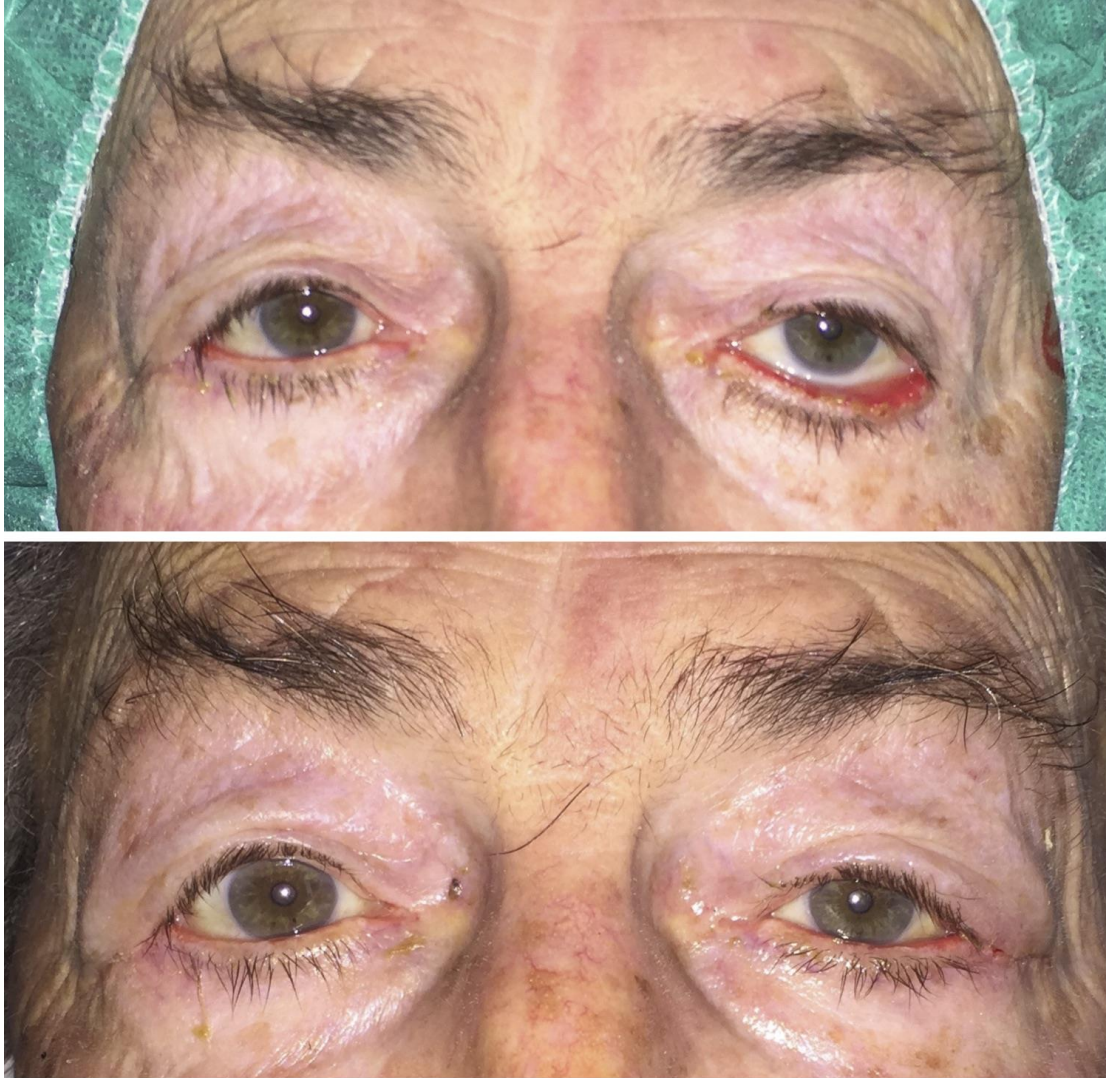


**Figure 6.** Lacrimal ectropion of the right lower eyelid (top) corrected with a Lazy-T procedure (bottom).





**Figure 7.** Left involutional lower lid ectropion. Top: Preoperative appearance.  
Bottom: postoperative (tarsal strip) result.





Um agradecimento muito especial à minha orientadora, a Doutora Sara Filipa Ribeiro Teixeira. Agradecimento também muito especial à família e aos amigos que sempre me acompanharam nesta jornada.

# **Anexos**

# Ophthalmic Plastic and Reconstructive Surgery

## INFORMATION FOR AUTHORS

***Ophthalmic Plastic and Reconstructive Surgery (OPRS)*** publishes Original Investigations describing clinical and laboratory investigations, Case Reports, Surgical Techniques, Anatomy & Physiology, and Reviews, and by invitation, Commentaries, Perspectives, and Editorials. The journal is owned and sponsored by the American Society of Ophthalmic Plastic and Reconstructive Surgery (ASOPRS) but welcomes submissions from authors who are not members of the Society.

### GENERAL INFORMATION AND POLICIES

*OPRS* subscribes to the policies outlined in the "Uniform Requirements for Manuscripts Submitted to Biomedical Journals" written by the International Committee of Medical Journal Editors ([www.ICMJE.org](http://www.ICMJE.org)).

**Authorship.** *OPRS* expects each author to have made a significant intellectual contribution to the design or execution of the project, to the writing of the manuscript, or both. Each author must take full responsibility for his or her contribution and must have approved the final manuscript. (Lundberg GD, Glass RM. What does authorship mean in a peer-reviewed medical journal? [editorial] *JAMA* 1996;276:75.)

**Papers from the ASOPRS Annual Scientific Symposia.** *OPRS* holds copyright and has the right of first refusal for all manuscripts derived from papers presented at the Annual Fall and Spring Symposia of the American Society of Ophthalmic Plastic and Reconstructive Surgery. Permission to submit such work elsewhere must be sought from the Editor. Presenters who violate this policy will be excluded by the Program Committee from participation in the Annual Scientific Symposia for 2 years.

**Prior and Repetitive Publication.** *OPRS* will not consider manuscripts that have appeared in part or in total in other publications, except in special circumstances by approval of the Editor. Updates of previously published studies that add little data to an existing publication will not be considered. Overlap between patient groups described in serial manuscripts must be acknowledged, and references to previous publications that include the same patients must be provided. Authors uncertain as to whether specific data represent prior or repetitive publication should alert the Editor in the transmittal letter and include copies of the publication(s) in question.

### Institutional Review Board Approval

For all manuscripts reporting data from studies involving human subjects, human-derived material, human medical records, or animals, formal ethical standards review and approval, or formal review and waiver, by an appropriate institutional review board (IRB) or ethics committee is required and must be described in the Methods section. Case reports presenting the clinical or surgical results on three patients or less, are generally not considered research, and therefore do not require IRB review. However, discussion or chart reviews of four or more patients, especially when informed consent is obtained, or any form of analysis or comparison with other studies is performed, may be considered research, and should be reviewed by an IRB board for approval or waiver. The IRB decision should be included in the introduction. For those investigators who do not have access to a formal institutional ethics review committee, a non-institutional regional IRB should be utilized. For non-USA investigators, who do not have access to any formal review board, please include a

statement to this effect in the Methods, and include a statement that the study adhered to the principles outlined in the Declaration of Helsinki (<http://www.wma.net/en/30publications/10policies/b3/>). Editors may request that authors provide documentation of the formal review and recommendation from the institutional review board or ethics committee responsible for oversight of the study.

### **Patient Consent**

For investigations on human subjects, state in the Methods section the manner in which informed consent was obtained from the study participants (ie, oral or written) and for US authors, that the study was HIPPA-compliant (<http://www.hhs.gov/ocr/privacy/index.html>).

### **Clinical Photographs**

All clinical and radiographic photographs that permit identification of the patient must have a signed consent by the patient or guardian, which is to be archived by the authors. This statement does not have to be submitted with the manuscript. A statement that this consent was obtained and is on file must be included in the Methods or Acknowledgement section. It is not acceptable to place bars over the patient's features, but in cases where permissions are unobtainable, the photographs must be very tightly cropped to the feature being displayed. If identification is still possible after cropping, *OPRS* cannot use the photograph. In the case of a patient who is deceased, written permission must be provided by the patient's next of kin.

**Animal Studies.** If animals were used in a study, the manuscript should include the appropriate IRB approval statement.

**Statistical Analysis.** *OPRS* strongly advises statistical consultation about data collection and analysis. Statistical methods must be identified in table footnotes, illustration legends, or text explanations. Software programs used for complex statistical analyses must be identified to enable reviewers to verify calculations.

## **MANUSCRIPT PREPARATION**

Manuscripts should be prepared using Microsoft Word. The manuscript should be paginated and double-spaced, with 1-inch margins and left-justified text. Please activate line numbering: File/pagesetup/page layout: Show line numbers, Line numbering: Continuous/Begin with line 1. Article length can be estimated according to the following formula: 1 journal page = 3 double-spaced type-written pages, OR 4 single-image figures, OR 2 tables. Submissions should include the intended article type as follows.

*Original Investigation:* These articles discuss original clinical or basic research with a detailed review of the literature that the research supports, expands or modifies. Submissions should be limited to six journal pages.

*Major Review:* These articles present a comprehensive review of the current literature summarizing our state of knowledge on a specific subject. Submissions are limited to 10 journal pages.

*Surgical Technique:* This article describes a new or modified surgical procedure, with or without case descriptions to illustrate the technique. Limited to 2 journal pages. Authors are encouraged to submit a video to accompany Surgical Technique articles to enhance the article content. Please refer to the Supplemental Digital Content section of this document for

instructions on submitting videos as Supplemental Digital Content, including directions on necessary file formats and required call-outs within the article.

*Anatomy & Physiology:* Includes original anatomical or physiological descriptions related to the eyelids, orbit, or lacrimal systems. Limited to 3 journal pages.

*Case Report:* These are clinical descriptions of one or more patients with a brief discussion of the relevant literature. Case reports will be published on-line only. Limited to 2 journal pages and 12 references. For all case reports originating in the USA, the introductory paragraph should state that collection and evaluation of protected patient health information was HIPAA-compliant.

*Images in Ophthalmic Plastic and Reconstructive Surgery:* Submissions should contain a maximum of 2 figures with accompanying figure legend. Video segments may also be included but submissions must include a high-resolution still image. Limited to ½ journal page. Please see [Images in OPRS](#) for more details.

*Letter to the Editor:* Letters are brief non-peer reviewed comments that relate to recently published papers in the journal; they may also present a brief discussion of the authors' clinical or research experience. Limited to 1 journal page. Letter titles should be "Re:" or "Reply re:" followed by the title of the article to be discussed, and the discussed article should be listed as a reference. If the letter pertains only to your own research experience, just an appropriate title is needed. Letters should be addressed to "To the Editor:".

**Title Page.** The title page should include the following information:

1. Title—no longer than 135 characters. Declarative titles should not be used.
2. Names of authors—provide first name, middle initial, last name, and advanced degrees or professional certification.
3. Institutional affiliation—indicate each author's affiliation during the course of the study in footnotes on the title page using superscript numbers, not symbols (e.g., John Doe<sup>1</sup>).
4. Meeting presentation—if the material has been presented previously, supply the name, place, and date of the meeting.
5. Financial support—identify all sources, public and private. Provide the agency name and city, company name and city, fellowship name, and grant number.
6. Proprietary interest statement—each author is expected to disclose any type of financial interest that is related to the manuscript, including stock or ownership of a business entity connected to a product described in the paper, paid consulting for the company or competing companies, or patent rights to a drug or piece of equipment. Authors must disclose personal or family ownership or potential rights to more than 1% of the company or competing company and whether they have any interest in marketing any product, drug, instrument, or piece of equipment discussed in the manuscript.
7. Running head—no longer than 60 characters.
8. Corresponding author—contact information for reprints.

**Précis.** On a separate page (page 2), include a one-sentence précis (35 words or fewer) summarizing the main finding or outcome of the study. The précis will appear under the title in the Table of Contents and should not duplicate the abstract conclusion.

**Structured Abstract.** Each manuscript must include a structured abstract of no more than 250 words (except for Case Reports; see below). The abstract must appear on a separate page (page 3) and should include four separate sections:

**Purpose:** Provide a concise statement of the study goal (e.g., the question to be answered or the hypothesis to be tested).

**Methods:** Identify the study design using a phrase such as randomized or nonrandomized clinical trial, case-controlled study, cross-sectional study, cohort study, case series, case report, meta-analysis, review, experimental study, or historical manuscript. Patient selection, interventions, and outcome measures must be defined. For studies involving human subjects, human-derived materials, or human medical records, include a statement declaring IRB approval, Helsinki adherence or HIPAA compliance as appropriate.

**Results:** Briefly summarize the principal measurements (data) obtained and relevant statistical analysis.

**Conclusions:** State the specific conclusions derived from the data analysis and their clinical significance.

**Abstract for Case Reports.** Provide an unstructured summary not to exceed 150 words on a separate page following the précis.

**Text.** All manuscripts must follow generally recognized standards for presenting scientific material. See the *CBE Manual for Authors, Editors, and Publishers*, 6th ed. (Council of Biology Editors; 1995). Type size (True Type fonts) should be 12 point. The **Introduction**, without a heading, should refer only to the most pertinent past publications and should not be an extensive review of the literature. **Methods** should be written with sufficient detail to permit others to duplicate the work. **Results** must be concise and not simply a reiteration of data presented in Tables. **Discussion** should be restricted to the significant findings presented. Digressions and speculation are not appropriate.

**References.** References should follow text and begin on a separate page. They must be double spaced and numbered consecutively in order of appearance in the text. References should be designated by superscript numbers following all punctuation (except semicolons and colons).

1. List only references that you have read and that are pertinent to the manuscript.
2. Cite only published studies as references. You may acknowledge "unpublished data" or "submitted" articles within parentheses in the text. Reference to a "personal communication" within parentheses in the text must be accompanied by a signed permission letter from the individual being cited.
3. Abstracts such as those published in the annual meeting programs of the American Society of Ophthalmic Plastic and Reconstructive Surgery (ASOPRS), the American Academy of Ophthalmology (AAO), or the Association for Research in Vision and Ophthalmology (ARVO) are considered "unpublished" and should be cited in parentheses in the text. For example: -by Smith et al. (*Invest Ophthalmol Vis Sci* 28 (Suppl):54, 1989).
4. Oral or poster presentations are similarly unpublished and may be cited only in parentheses in the text. Platform and poster presentations at annual meetings are customarily indicated: (Smith AB, presented at the AAO Annual Meeting, Atlanta, 1995).
5. Books or articles "in press" may be cited as numbered references. Such citations should be updated before publication, if possible.

Journal abbreviations should be those used by the National Library of Medicine, as found in *Index Medicus*. If in doubt as to the correct abbreviation, cite the complete journal name.

Do not underline journal titles, and do not use periods in abbreviations of journal titles or in author initials. Please follow precisely the format and punctuation shown in the following examples.

**Journal Article–(If four or fewer authors, list all)** Anderson RL, Beard C. The levator aponeurosis: attachments and their clinical significance. *Arch Ophthalmol* 1977;95:1437– 41.

**Journal Article–(If five or more authors, list only the first three and add et al)** Meyer DR, Bui HX, Carlson JA, et al. Silicon granulomas and dermatomyositis-like changes associated with chronic eyelid edema after silicone breast implant. *Ophthal Plast Reconstr Surg* 1998;14:182–8.

**Chapter in a Book**–Kaltreider SA, Sherman DD, McGetrick JJ. Eyelid trauma. In: Dortzbach RK, ed. *Ophthalmic plastic surgery: prevention and management of complications*. New York: Raven Press, 1994:157–74.

**Book**–Miller NR. *Walsh and Hoyt’s clinical neuro-ophthalmology*. 4th ed. Vol. 4. Baltimore: Williams & Wilkins; 1991; 2102–14.

**Letter to the Editor**–Sneed SR, Blodi CF, Berger BB, et al. Pneumocystis carinii choroiditis in patients receiving inhaled pentamidine [letter]. *N Engl J Med* 1990;322:936 –7.

**Online Journal Article**–LaPorte RE, Marler E, Akazawa S, Sauer F. The death of biomedical journals. *JAMA* [serial online]. 1996;310:1387–90. Available at <http://www.jama.com/jama/archive/6991ed2.htm>. Accessed June 16, 1997.

**Web Site**–Health Care Financing Administration. 1996 statistics at a glance. Available at: <http://www.hcfa.gov/stats/stathili.htm>. Accessed December 2, 1997.

**Tables.** Do not embed tables within the body of the manuscript. Each table must be numbered consecutively using Arabic numbers, be mentioned in the text, and be titled. Each column must have a heading. All abbreviations must be explained in the legend. Please do not place more than one table on a page.

**Legends.** Figure legends (photos, drawings, graphs) should be placed at the end of the manuscript. Do not embed figures within the body of the manuscript. Figures must be numbered consecutively as they appear in the text. For histologic figures, stains and magnifications should be noted in the legend. Any figure that has been published elsewhere should have an acknowledgment to the original source; a copy of the release to publish the figure, signed by the copyright holder, must also be submitted. Legends must identify all symbols or letters that appear on the prints.

**Abbreviations.** Restrict abbreviations to those that are widely used and understood. Avoid abbreviations that have meaning only in the context of your specific manuscript. An abbreviation should appear first in parentheses immediately after the term or phrase to which it refers. Abbreviations may only be used if the term appears five or more times in the text. Abbreviations are not used in abstracts.

**Drug/Manufacturer Names.** Use generic names only in the text body. Include the trade name of a particular drug and the manufacturer’s name and location, cited in parentheses,

after the first use of the generic name. In the case of equipment, include manufacturer's name, city, state, and/or country.

**Illustrations.** Illustrations should be prepared according to the image guidelines for online manuscript submission.

**Copyright.** Consideration of manuscripts for publication in *Ophthalmic Plastic and Reconstructive Surgery* is dependent on the assurance that the material (in whole or part) is not under consideration by another journal, is not in press in any other format, and has not been previously published. Each author must sign a statement transferring copyright ownership to the American Society of Ophthalmic Plastic and Reconstructive Surgery. Copyright transfer forms are printed in each issue of the journal and are available online at the journal's Web site ([www.op-rs.com](http://www.op-rs.com)).

**Conflicts of Interest.** Authors must state all possible conflicts of interest in the manuscript, including financial, consultant, institutional and other relationships that might lead to bias or a conflict of interest. If there is no conflict of interest, this should also be explicitly stated as none declared. All sources of funding should be acknowledged in the manuscript. All relevant conflicts of interest and sources of funding should be included on the title page of the manuscript with the heading "Conflicts of Interest and Source of Funding:".

For example: Conflicts of Interest and Source of Funding: A has received honoraria from Company Z. B is currently receiving a grant (#12345) from Organization Y, and is on the speaker's bureau for Organization X – the CME organizers for Company A. For the remaining authors none were declared.

Each author must complete and submit the journal's copyright transfer agreement, which includes a section on the disclosure of potential conflicts of interest based on the recommendations of the International Committee of Medical Journal Editors, "Uniform Requirements for Manuscripts Submitted to Biomedical Journals" ([www.icmje.org/update.html](http://www.icmje.org/update.html)).

A copy of the form is made available to the submitting author within the Editorial Manager submission process. Co-authors will automatically receive an Email with instructions on completing the form upon submission.

### **Open access**

LWW's hybrid open access option is offered to authors whose articles have been accepted for publication. With this choice, articles are made freely available online immediately upon publication. Authors may take advantage of the open access option at the point of acceptance to ensure that this choice has no influence on the peer review and acceptance process. These articles are subject to the journal's standard peer-review process and will be accepted or rejected based on their own merit.

Authors of accepted peer-reviewed articles have the choice to pay a fee to allow perpetual unrestricted online access to their published article to readers globally, immediately upon publication. The article processing charge for *Ophthalmic Plastic and Reconstructive Surgery* is \$2,100. The article processing charge for authors funded by the Research Councils UK (RCUK) is \$2,640. The publication fee is charged on acceptance of the article and should be paid within 30 days by credit card by the author, funding agency or institution. Payment



must be received in full for the article to be published open access. Any additional standard publication charges, such as for color images, will also apply.

- ***Authors retain copyright***

Authors retain their copyright for all articles they opt to publish open access. Authors grant LWW a license to publish the article and identify itself as the original publisher.

- ***Creative Commons license***

Articles opting for open access will be freely available to read, download and share from the time of publication. Articles are published under the terms of the Creative Commons License Attribution-NonCommercial No Derivative 3.0 which allows readers to disseminate and reuse the article, as well as share and reuse of the scientific material. It does not permit commercial exploitation or the creation of derivative works without specific permission. To view a copy of this license visit: <http://creativecommons.org/licenses/by-nc-nd/3.0>.

- ***Compliance with NIH, RCUK, Wellcome Trust and other research funding agency accessibility requirements***

A number of research funding agencies now require or request authors to submit the post-print (the article after peer review and acceptance but not the final published article) to a repository that is accessible online by all without charge. As a service to our authors, LWW identifies to the National Library of Medicine (NLM) articles that require deposit and transmits the post-print of an article based on research funded in whole or in part by the National Institutes of Health, Howard Hughes Medical Institute, or other funding agencies to PubMed Central. The revised Copyright Transfer Agreement provides the mechanism. LWW ensures that authors can fully comply with the public access requirements of major funding bodies worldwide. Additionally, all authors who choose the open access option will have their final published article deposited into PubMed Central.

RCUK and Wellcome funded authors can choose to publish their paper as open access with the payment of an article process charge (gold route), or opt for their accepted manuscript to be deposited (green route) into PMC with an embargo.

With both the gold and green open access options, the author will continue to sign the Copyright Transfer Agreement (CTA) as it provides the mechanism for LWW to ensure that the author is fully compliant with the requirements. After signature of the CTA, the author will then sign a License to Publish where they will then own the copyright. Those authors who wish to publish their article via the gold route will be able to publish under the terms of the Attribution 3.0 (CCBY) License. To view of a copy of this license visit: <http://creativecommons.org/licenses/by/2.0/>. Those authors who wish to publish their article via the green route will be able to publish under the rights of the Attribution Non-commercial 3.0 (CCBY NC) license (<http://creativecommons.org/licenses/by-nc/2.0/>).

It is the responsibility of the author to inform the Editorial Office and/or LWW that they have RCUK funding. LWW will not be held responsible for retroactive deposits to PMC if the author has not completed the proper forms.

**FAQ for open access**

<http://links.lww.com/LWW-ES/A48>

**Cancer Classification Scheme.** Authors should use the American Joint Commission on Cancer classification scheme when describing patients with ophthalmic malignancies; see American Joint Committee on Cancer. *AJCC Cancer Staging Manual*, 7th Edition, New York, Springer, 2010.

**Corresponding Author.** The editorial office must be supplied with phone and fax numbers, and the e-mail address for the corresponding author. Please do not choose a corresponding author who will not be available to respond to editorial queries during the evaluation and publication process. The editorial office must be notified of any changes in the order of authorship, author name change, or address or phone number of the corresponding author. Always indicate the manuscript number in subsequent communications or correspondence.

**Receipt of Manuscript, Review Process, and Revisions.** Each manuscript will be acknowledged via e-mail in the order received. The acknowledgment will note the number assigned to the manuscript; this number should be referenced during all subsequent communications about the manuscript. *OPRS* does not reveal the identity of its reviewers but does provide pertinent comments to the corresponding author. Usually two and sometimes several reviewers will participate in the review of a manuscript. Re-review may be required after revision if, in the judgment of the Editor, sufficient modification of the manuscript or data justifies another review cycle.

**ONLINE MANUSCRIPT SUBMISSION**

*Ophthalmic Plastic and Reconstructive Surgery* accepts online submission of manuscripts through Editorial Manager™, which is linked to [www.op-rs.com](http://www.op-rs.com). The site contains instructions and advice on how to use the system, guidance on the creation/scanning and saving of electronic art, and supporting documentation.

Although Editorial Manager presently accepts many file formats, authors are advised to use only those that are acceptable to Lippincott Williams & Wilkins, the publisher, in order to ensure proper publication in the print issues. Please see individual sections below for specific file requirements for text, tables, and figures. Please review the files as they upload to ensure each file name has a corresponding file extension (i.e., .doc, .tif). Adherence to the guidelines is essential if efficient and expeditious processing of your manuscript is to be achieved. Manuscripts not submitted in the correct format will be returned to authors for revision before peer review. Authors who submit their manuscripts through Editorial Manager are asked not to send e-mailed or hard copies of the manuscript to the Journal's editorial office.

**File Formats.** Following the guidelines for manuscript preparation, text and tables should be prepared using Microsoft Word. Figures should be submitted in TIFF format.

**File Size.** Manuscripts are distributed to reviewers via the Web. However, reviewers who use telephone modems may experience unacceptable download delays if the files are too large. Some simple measures can avoid unnecessarily large files. Do not scan pages of text. Do not scan printed figures unless no original digital or film image exists. If a scanned figure is unavoidable, please use Adobe PhotoShop or a similar program to reduce the file size (not

necessarily the image size). For example, crop the picture to exclude surrounding "white space." Black-and-white line drawings or grayscale figures should not be saved as color documents; this will increase file size without increasing the information content of the file.

## **SUPPLEMENTAL DIGITAL CONTENT**

**Supplemental Digital Content (SDC).** Authors may submit SDC that enhances their article's text via Editorial Manager to LWW journals to be considered for online posting. SDC may include standard media such as text documents, graphs, audio, video, etc. On the Attach Files page of the submission process, please select Supplemental Audio, Video, or Data for your uploaded file as the Submission Item. If an article with SDC is accepted, our production staff will create a URL with the SDC file. The URL will be placed in the call-out within the article. SDC files are not copyedited by LWW staff; they will be presented digitally as submitted. For a list of all available file types and detailed instructions, please visit <http://links.lww.com/A142>.

**SDC Call-outs.** SDC must be cited consecutively in the text of the submitted manuscript. Citations should include the type of material submitted (Audio, Figure, Table, etc.), be clearly labeled as "Supplemental Digital Content," include the sequential list number, and provide a description of the supplemental content. All descriptive text should be included in the call-out, as it will not appear elsewhere in the article. Example: We performed many tests on the degrees of flexibility in the elbow (see Video, Supplemental Digital Content 1, which demonstrates elbow flexibility) and found our results inconclusive.

**List of SDC.** A listing of SDC must be submitted at the end of the manuscript file. Include the SDC number and file type of the SDC. This text will be removed by our production staff and not be published. Example: Supplemental Digital Content 1.wmv.

**SDC File Requirements.** All acceptable file types are permissible up to 10 MBs. For audio or video files greater than 10 MBs, authors should first query the journal office for approval. For a list of all available file types and detailed instructions, please visit <http://links.lww.com/A142>.

## **IMAGES**

Photomicrographs of histopathologic sections must be submitted in color. Authors are encouraged to submit other figures in color, when appropriate. The cost to authors for color reproduction is \$100 per figure. A figure can comprise one or more images/figure parts. Color costs can be reduced by judicious grouping of color images into composite figures. If figures are submitted in color, it is assumed the author intends to pay for color reproduction.

Do not paste figures into word processing documents; submit them as separate files, without their captions. Label each file with its figure number and upload the figures in numerical order.

The following summarizes our printer's guidelines for image preparation. Additional details can be found at <http://cpc.cadmus.com/da/guidelines.asp>. Please follow these guidelines carefully. Image manipulation by the printer will substantially increase the cost to authors.

**Please note:** Images should be submitted in TIFF format. JPEG, GIFF, PowerPoint, Excel, CorelDRAW, Quattro Pro, MS Word, and downloaded Internet image files are not acceptable at this time.

**Resolution:** The minimum requirements for resolution are:

- 1200 DPI/PPI for monochrome.
- For purely black and white images, such as line graphs and artistic renderings.
- 300 DPI/PPI for halftones (CMYK/grayscale).

For color or black-and-white images containing pictures only, including photographs not containing text labeling or thin lines.

- 600 DPI/PPI for combination halftones (CMYK/grayscale)

For color or black-and-white images containing pictures and text labeling and/or thin lines.

Lower resolutions may compromise print quality; higher resolutions will not improve output quality and will only increase file size.

**Color Mode.** Color images must be submitted in CMYK (cyan, magenta, yellow, black) mode, **not** RGB (red, green, blue) mode. Radiographs should be submitted in grayscale mode. Black-and-white line art can be submitted in grayscale or bitmap mode.

**Cropping and sizing.** All graphics should be submitted at their actual size; that is, they should be 100% of their print dimensions so that no scaling is necessary. Images should fit a column width of 3.5 inches. Crop figures (or change the pagesize of your document) so that no unnecessary white space is left bordering the figure.

**Line Art.** Line art must be submitted as a TIFF file at a resolution of 1200 dpi. If prepared in PowerPoint or Excel, line art can be imported into Photoshop and converted to TIFF format or can be printed on a photoquality printer and then rescanned at 1200 dpi.